

Form PTO-1449 (modified)

List of Patents and Publications for Applicant's

## INFORMATION DISCLOSURE STATEMENT

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Atty. Docket No.

GOUD:023USD1

Serial No.

10/664,603

Applicant

Guy A. Rouleau *et al.*

Filing Date:

September 17, 2003

Group:

Unknown

U.S. Patent Documents

See Page 1

Foreign Patent Documents

See Page 1

Other Art

See Page 1

## U.S. Patent Documents

Exam. Init.	Ref. Des.	Document Number	Date	Name	Class	Sub Class	Filing Date of App.
SL	A1	5,223,409	6-29-93	Ladner et al.	435	69.7	3-1-91

## Foreign Patent Documents

Exam. Init.	Ref. Des.	Document Number	Date	Country	Class	Sub Class	Translation Yes/No
SL	B1	WO 99/21875	5-6-99	PCT			

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Exam. Init.	Ref. Des.	Citation
SL	C1	Andermann, E., Genetic Basis of the Epilepsies, Raven Press, New York, pp. 355-374, 1982.
	C2	Anderson et al., "Use of cyclosporin A in establishing Epstein-Barr virus-transformed human lymphoblastoid cell lines," <i>In Vitro</i> , 20:856-858, 1984.
	C3	Annegers et al., Genetic Basis of the Epilepsies, Raven Press, New York, pp.151-159, 1982.
	C4	Baker et al., "Cell proliferation kinetics of normal and tumor tissue in vitro: quiescent reproductive cells and the cycling reproductive fraction," <i>Cell Prolif.</i> , 28:1-15, 1995.
	C5	Barker et al., "GABA actions on the excitability of cultured CNS neurons," <i>Neurosci. Lett.</i> , 47:313-318, 1984.
	C6	Bar-Sagi et al., "Negative modulation of sodium channels in cultured chick muscle cells by the channel activator batrachotoxin," <i>J. Biol. Chem.</i> , 260:4740-4744, 1985.
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	C10	Biervet et al., "A potassium channel mutation in neonatal human epilepsy," <i>Science</i> , 279:403-406, 1998.

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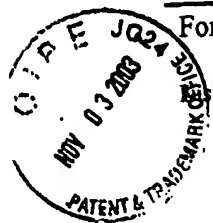
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U.S. Patent Documents

See Page 1

Foreign Patent Documents

See Page 1

Other Art

See Page 1

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	C12	Cardell et al., <i>Agnew. Chem. Int. Ed. Engl.</i> , 33:2061-2063, 1994.
	C13	Charlier et al., "A pore mutation in a novel KQT-like potassium channel gene in an idiopathic epilepsy family," <i>Nat. Genet.</i> , 18:53-55, 1998.
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	C16	Cho et al., "An Unnatural Biopolymer," <i>Science</i> , 261:1303-1305, 1993.
	C17	Clare et al., "Voltage-gated sodium channels as therapeutic targets," <i>Drug Discovery Today</i> , 5:506-520, 2000.
	C18	Corey et al., "The occurrence of epilepsy and febrile seizures in Virginian and Norwegian twins," <i>Neurology</i> , 41:1433-1436, 1991.
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	C21	DeWitt et al., "'Diversomers': an approach to nonpeptide, nonoligomeric chemical diversity," <i>Proc. Natl. Acad. Sci. USA</i> , 90:6909-6913, 1993.
	C22	Elliot et al., "Bin1 functionally interacts with Myc and inhibits cell proliferation via multiple mechanisms," <i>Oncogene</i> , 18:3564-3573, 1999.
	C23	Elmslie et al., "Genetic mapping of a major susceptibility locus for juvenile myoclonic epilepsy on chromosome 15q," <i>Hum. Mol. Genet.</i> , 6:1329-1334, 1997.
↓	C24	Engel et al., <i>Epilepsy: A Comprehensive Textbook</i> , Lippincott-Raven, Philadelphia, 1-7 (1), 1997.

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Guy A. Rouleau *et al.*Filing Date:  
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UnknownU.S. Patent Documents  
See Page 1Foreign Patent Documents  
See Page 1Other Art  
See Page 1

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	C27	Fodor et al., "Multiplexed biochemical assays with biological chips," <i>Nature</i> , 364:555-556, 1993.
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	C31	Greenberg et al., "Juvenile myoclonic epilepsy (JME) may be linked to the BF and HLA loci on human chromosome 6," <i>Am. J. Med. Genet.</i> , 31:185-192, 1988.
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U.S. Patent Documents

See Page 1

Foreign Patent Documents

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	C42	Leppert et al., "Benign familial neonatal convulsions linked to genetic markers on chromosome 20," <i>Nature</i> , 337:647-648, 1989.
	C43	Lewis et al., "Genetic heterogeneity in benign familial neonatal convulsions: identification of a new locus on chromosome 8q," <i>Am. J. Hum. Genet.</i> , 53:670-675, 1993.
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	C51	Moulard et al., "Identification of a new locus for generalized epilepsy with febrile seizures plus (GEFS+) on chromosome 2q24-q33," <i>Am. J. Hum. Genet.</i> , 65:1396-1400, 1999.

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Unknown

U.S. Patent Documents

See Page 1

Foreign Patent Documents

See Page 1

Other Art

See Page 1

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SL	C52	Muir et al., "Phase II clinical trial of sipatrigine (619C89) by continuous infusion in acute stroke," <i>Cerebrovascular Diseases</i> , 10:431-436, 2000.
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	C60	Pugsley et al., "Effects of bisaramil, a novel class I antiarrhythmic agent, on heart, skeletal muscle and brain Na <sup>+</sup> channels," <i>Eur. J. Pharmacol.</i> , 342:93-104, 1998.
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	C63	Sillampää et al., "Genetic factors in epileptic seizures: evidence from a large twin population," <i>Acta Neurol. Scand.</i> , 84:523, 1991.
	C64	Singh et al., "A novel potassium channel gene, KCNQ2, is mutated in an inherited epilepsy of newborns," <i>Nat. Genet.</i> , 18:25-29, 1998.
✓	C65	Sjolander et al., "Integrated fluid handling system for biomolecular interaction analysis," <i>Anal. Chem.</i> , 63:2338-2345, 1991.

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September 17, 2003Group:  
Unkn. wnU.S. Patent Documents  
*See Page 1*Foreign Patent Documents  
*See Page 1*Other Art  
*See Page 1*

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	C70	Wallace et al., "Febrile seizures and generalized epilepsy associated with a mutation in the Na <sup>+</sup> -channel beta1 subunit gene SCN1B," <i>Nature Genet.</i> , 19:366-370, 1998.
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## INFORMATION DISCLOSURE STATEMENT

(Use several sheets if necessary)

Filing Date:

September 17, 2003

Group:

1632

U.S. Patent Documents

See Page 1

Foreign Patent Documents

See Page 1

Other Art

See Page 1

## U.S. Patent Documents

Exam. Init.	Ref. Des.	Document Number	Date	Name	Class	Sub Class	Filing Date of App.

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		Filing Date: September 17, 2003	Group: 1639
U.S. Patent Documents <i>See Page 1</i>	Foreign Patent Documents <i>See Page 1</i>	Other Art <i>See Page 1</i>	

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